

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1 1. (Original) A method for determining a PCR amplified nucleic acid, said
2 method comprising:
 - 3 a) contacting a sample comprising nucleic acid to be PCR amplified with at
4 least one polymer comprising at least one linked energy donor moiety and at least one linked
5 energy acceptor moiety wherein said donor and acceptor moieties are separated by at least a
6 portion of a probing nucleobase sequence and wherein said polymer does not form a stem and
7 loop hairpin and is further characterized in that the efficiency of transfer of energy between said
8 donor and acceptor moieties, when the polymer is solvated in aqueous solution, is substantially
9 independent of at least two variables selected from the group consisting of:
 - 10 i) nucleobase sequence length separating the at least one energy
11 donor moiety from the at least one energy acceptor moiety;
 - 12 ii) spectral overlap of the at least one linked energy donor moiety and
13 the at least one linked energy acceptor moiety;
 - 14 iii) presence or absence of magnesium in the aqueous solution; and the
15 iv) ionic strength of the aqueous solution;
 - 16 b) contacting the sample with primers and other reagents for PCR
17 amplification;
 - 18 c) PCR amplifying the nucleic acid;
 - 19 d) determining hybridization of the polymer to a target sequence within the
20 amplified nucleic acid wherein the target sequence present in the sample is correlated with a
21 change in detectable signal associated with at least one donor or acceptor moiety of the polymer;
22 and

23 e) determining the presence, absence or amount of PCR amplified nucleic
24 acid in the sample.

1 2. (Original) The method of claim 1, wherein the nucleic acid to be
2 amplified is contained within a plasmid.

1 3. (Original) The method of claim 1, wherein the PCR amplification is
2 traditional.

1 4. (Original) The method of claim 1, wherein the PCR amplification is
2 asymmetric.

1 5. (Original) The method of claim 1, wherein the PCR amplification is
2 performed as a closed tube (homogeneous) assay.

1 6. (Original) The method of claim 1, wherein the amplified nucleic acid is
2 quantitated.

1 7. (Original) The method of claim 1, wherein the presence or absence of the
2 amplified nucleic acid is determined.

1 8. (Original) The method of claim 1, wherein the polymer is a peptide
2 nucleic acid.

1 9. (Original) A method for determining a PCR amplified nucleic acid, said
2 method comprising:

3 a) contacting a sample comprising nucleic acid to be PCR amplified with a
4 polymer comprising:

5 i) a probing nucleobase sequence for probing a target sequence to
6 which the probing nucleobase sequence is complementary or substantially
7 complementary;

8 ii) at least one energy donor moiety that is linked to the probing
9 nucleobase sequence; and
10 iii) at least one energy acceptor moiety that is linked to the probing
11 nucleobase sequence wherein the at least one donor moiety is separated from the at least
12 one acceptor moiety by at least a portion of the probing nucleobase sequence;
13 b) contacting the sample with primers and other reagents for PCR
14 amplification;
15 c) PCR amplifying the nucleic acid;
16 d) determining hybridization of the polymer to a target sequence within the
17 amplified nucleic acid wherein the target sequence present in the sample is correlated with a
18 change in detectable signal associated with at least one donor or acceptor moiety of the polymer;
19 and
20 e) determining the presence, absence or amount of PCR amplified nucleic
21 acid in the sample.

1 10. (Original) The method of claim 9, wherein the nucleic acid to be
2 amplified is contained within a plasmid.

1 11. (Original) The method of claim 9, wherein the PCR amplification is
2 traditional.

1 12. (Original) The method of claim 9, wherein the PCR amplification is
2 asymmetric.

1 13. (Original) The method of claim 9, wherein the PCR amplification is
2 performed as a closed tube (homogeneous) assay.

1 14. (Original) The method of claim 9, wherein the amplified nucleic acid is
2 quantitated.

1 15. (Original) The method of claim 9, wherein the presence or absence of the
2 amplified nucleic acid is determined.

1 16. (Original) The method of claim 9, wherein the polymer is a peptide
2 nucleic acid.